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NEWS 1		Web Page for STN Seminar Schedule - N. America
NEWS 2	AUG 06	CAS REGISTRY enhanced with new experimental property tags
NEWS 3	AUG 06	FSTA enhanced with new thesaurus edition
NEWS 4	AUG 13	CA/CAplus enhanced with additional kind codes for granted patents
NEWS 5	AUG 20	CA/CAplus enhanced with CAS indexing in pre-1907 records
NEWS 6	AUG 27	Full-text patent databases enhanced with predefined patent family display formats from INPADOCDB
NEWS 7	AUG 27	USPATOLD now available on STN
NEWS 8	AUG 28	CAS REGISTRY enhanced with additional experimental spectral property data
NEWS 9	SEP 07	STN AnaVist, Version 2.0, now available with Derwent World Patents Index
NEWS 10	SEP 13	FORIS renamed to SOFIS
NEWS 11	SEP 13	INPADOCDB enhanced with monthly SDI frequency
NEWS 12	SEP 17	CA/CAplus enhanced with printed CA page images from 1967-1998
NEWS 13	SEP 17	CAplus coverage extended to include traditional medicine patents
NEWS 14	SEP 24	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS 15	OCT 02	CA/CAplus enhanced with pre-1907 records from Chemisches Zentralblatt
NEWS 16	OCT 19	BEILSTEIN updated with new compounds
NEWS 17	NOV 15	Derwent Indian patent publication number format enhanced
NEWS 18	NOV 19	WPIX enhanced with XML display format
NEWS 19	NOV 30	ICSD reloaded with enhancements
NEWS 20	DEC 04	LINPADOCDB now available on STN
NEWS 21	DEC 14	BEILSTEIN pricing structure to change
NEWS 22	DEC 17	USPATOLD added to additional database clusters
NEWS 23	DEC 17	IMSDRUGCONF removed from database clusters and STN
NEWS 24	DEC 17	DGENE now includes more than 10 million sequences
NEWS 25	DEC 17	TOXCENTER enhanced with 2008 MeSH vocabulary in MEDLINE segment
NEWS 26	DEC 17	MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
NEWS 27	DEC 17	CA/CAplus enhanced with new custom IPC display formats

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 NEWS 28  DEC 17  STN Viewer enhanced with full-text patent content
                  from USPATOLD
 NEWS 29  JAN 02  STN pricing information for 2008 now available

 NEWS EXPRESS  19 SEPTEMBER 2007:  CURRENT WINDOWS VERSION IS V8.2,
               CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
               AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.

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FILE 'HOME' ENTERED AT 18:12:09 ON 15 JAN 2008

=> file reg
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                     0.21       0.21

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DICTIONARY FILE UPDATES:  14 JAN 2008  HIGHEST RN 960583-85-1

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http://www.cas.org/support/stngen/stndoc/properties.html

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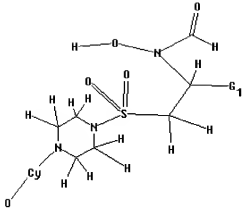

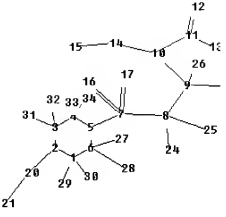

chain nodes :
7 8 9 10 11 12 13 14 15 16 17 19 20 21 23 24 25 26 27 28 29
30 31 32 33 34
ring nodes :
1 2 3 4 5 6
chain bonds :
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8-24 8-25 9-10 9-19 9-26 10-11 10-14 11-12 11-13 14-15 20-21 21-23
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6
exact/norm bonds :
1-2 1-6 2-3 2-20 3-4 4-5 5-6 5-7 7-8 7-16 7-17 9-10 9-19 10-11 10-14
11-12 20-21 21-23
exact bonds :
1-29 1-30 3-31 3-32 4-33 4-34 6-27 6-28 8-9 8-24 8-25 9-26 11-13 14-15

isolated ring systems :
containing 1 :

G1:Cy,Ak

G2:Cb,Ak

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:CLASS
11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 19:CLASS
20:Atom 21:CLASS
23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS 29:CLASS 30:CLASS
31:CLASS 32:CLASS
33:CLASS 34:CLASS

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L1      STRUCTURE UPLOADED

=> s l1 fulll
COMBINATION OF STRUCTURE AND TEXT TERMS NOT VALID
The query entered contains both search terms created by
structure-building or screen commands and text search terms.  L#s
created via the STRUCTURE or SCREEN commands must be searched in the
structures files separately from text terms or profiles.  The L#
answer sets from structure searches can be used in crossover searches
and can be combined with text terms.

=> s l1 full
FULL SEARCH INITIATED 18:13:51 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -      837 TO ITERATE

100.0% PROCESSED      837 ITERATIONS                              51 ANSWERS
SEARCH TIME: 00.00.01

L2           51 SEA SSS FUL L1

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COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                   178.82     179.03

FILE 'CAPLUS' ENTERED AT 18:13:55 ON 15 JAN 2008
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FILE LAST UPDATED: 14 Jan 2008  (20080114/ED)

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=> s l2 full
L3            3 L2

=> d ibib abs hitstr tot

L3   ANSWER 1 OF 3  CAPLUS  COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:         2006:630134  CAPLUS  Full-text
DOCUMENT NUMBER:          145:103718
TITLE:                    Preparation of (piperazinylsulfonylmethyl)alkynyl
                          hydroxamates and analogs as matrix metalloprotease
```

inhibitors and medical uses thereof

INVENTOR(S): Swinnen, Dominique; Bombrun, Agnes; Gerber, Patrick; Jorand-Lebrun, Catherine

PATENT ASSIGNEE(S): Applied Research Systems Ars Holding N.V., Neth. Antilles

SOURCE: PCT Int. Appl., 151 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006067114	A1	20060629	WO 2005-EP56910	20051219
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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
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CA 2589367	A1	20060629	CA 2005-2589367	20051219
EP 1828160	A1	20070905	EP 2005-826371	20051219
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KR 2007095950	A	20071001	KR 2007-716256	20070716
PRIORITY APPLN. INFO.:			EP 2004-106814 A	20041221
			US 2004-638257P P	20041222
			WO 2005-EP56910 W	20051219

OTHER SOURCE(S): MARPAT 145:103718

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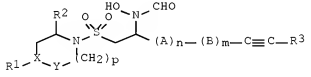

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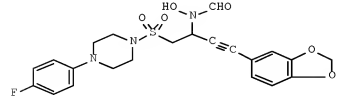

II

AB Title compds. I [wherein A, B = (un)substituted CH2; R1 = (hetero)aryl or (hetero)cycloalkyl; R2 = H, alkyl, alkenyl or alkynyl; R3 = H, alkyl, (hetero)aryl, etc.; X = C, CH or N; Y = CH, CH2, -C=CH-, etc.; m = 0-2, n = 0-1; p = 1-2] and stereoisomers or pharmaceutically acceptable salts thereof were prepared as matrix metalloprotease (MMP) inhibitors. Some related intermediates were claimed. For instance, successive lithiation of 1-(4-fluorophenyl)-4-(methylsulfonyl)piperazine with lithium bis(trimethylsilyl)amide, reaction with di-Et chlorophosphate, olefination with 3-(1,3-Benzodioxol-5-yl)-2-propynal (69% yield for three steps), nucleophilic addition of the resultant α,β-unsatd. sulfone with hydroxylamine (81% yield), and N-formylation with formic acetic anhydride generated in situ from acetic anhydride and formic acid (50% yield) gave hydroxamate II. This product showed inhibition against MMP-1 and MMP-12 with IC50 values of > 5000 nM and 46 nM, resp. Other biol. activities were also disclosed. Therefore, I and their pharmaceutical comps. are useful for the treatment and/or prophylaxis of autoimmune disorders, inflammatory diseases, cardiovascular diseases, neurodegenerative diseases, cancer, respiratory diseases and fibrosis.

IT 895573-30-5P 895573-54-3P 895573-58-7P
895573-63-4P 895573-92-9P 895573-94-1P
895574-00-2P 895574-01-3P 895574-08-0P
895574-16-0P 895574-21-7P 895574-24-0P
895574-30-8P 895574-31-9P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of (piperazinylsulfonylmethyl)alkynyl hydroxamates and analogs as matrix metalloprotease inhibitors and medical uses thereof)

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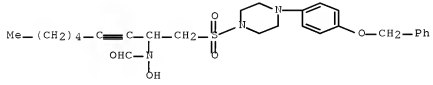

RN 895573-54-3 CAPLUS
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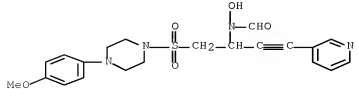

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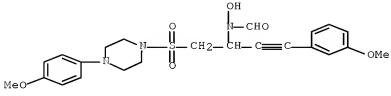

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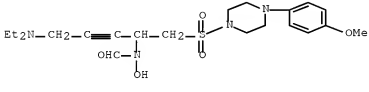

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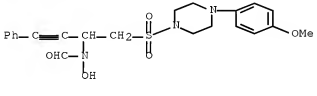

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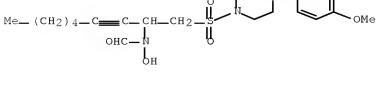

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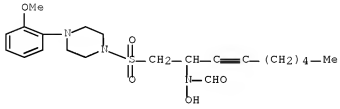

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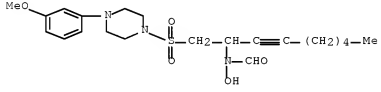

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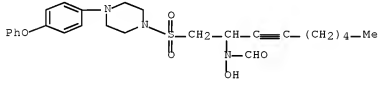

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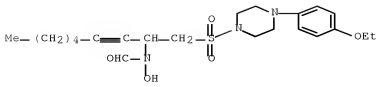

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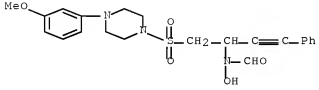

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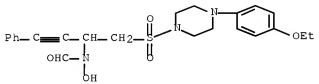

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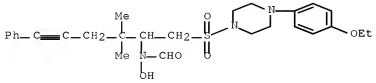

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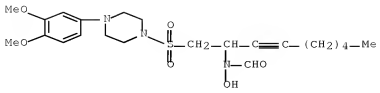

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:14380 CAPLUS Full-text
DOCUMENT NUMBER: 142:114099
TITLE: Preparation of N-[(4-substituted piperazine-1-sulfonylmethyl)alkyl]-N-hydroxyformamides as metalloproteinase inhibitors
INVENTOR(S): Finlay, Maurice Raymond Verschoyle; Waterson, David
PATENT ASSIGNEE(S): AstraZeneca AB, Swed.; AstraZeneca UK Limited
SOURCE: PCT Int. Appl., 57 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005000822	A1	20050106	WO 2004-GB2702	20040623
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004251104	A1	20050106	AU 2004-251104	20040623
CA 2529468	A1	20050106	CA 2004-2529468	20040623
EP 1644340	A1	20060412	EP 2004-743053	20040623
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CN 1812974	A	20060802	CN 2004-80018244	20040623
BR 2004011929	A	20060815	BR 2004-11929	20040623
JP 2007516164	T	20070621	JP 2006-516467	20040623
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NO 2006000444	A	20060322	NO 2006-444	20060127

PRIORITY APPLN. INFO.: SE 2003-1922 A 20030627
WO 2004-GB2702 W 20040623

OTHER SOURCE(S): MARPAT 142:114099
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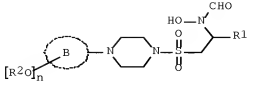

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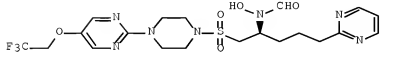


II

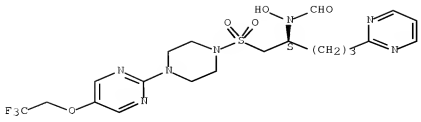
AB The title compds. I [ring B = monocyclic aryl ring having six ring atoms or a monocyclic heteroaryl ring having up to six ring atoms and containing one or more ring heteroatoms wherein each said heteroatom is nitrogen; R2 = alkyl or aryl, which said group is substituted by one or more fluorine groups; n = 1-3; R1 = (un)substituted alkyl, cycloalkyl, heterocycloalkyl, aryl, heteroaryl, alkyl-aryl, alkyl-heteroaryl, alkyl-cycloalkyl or alkyl-heterocycloalkyl], useful in the treatment of a disease condition mediated by one or more metalloproteinase enzymes, were prepared E.g., a multi-step synthesis of (S)-II, starting from 5-iodo-2-[4- (methylsulfonyl)piperazin-1-yl]pyrimidine, was given. In general, the compds. I demonstrate IC50 values in the range of 0.01 to 1000 nM against collagenase 3. The pharmaceutical composition comprising the compound I is disclosed.

IT 823197-00-8P 823197-01-9P 823197-02-0P
823197-03-1P 823197-04-2P 823197-05-3P
823197-06-4P 823197-07-5P 823197-08-6P
823197-09-7P 823197-10-0P 823197-11-1P
823197-12-2P 823197-13-3P 823197-14-4P
823197-15-5P 823197-16-6P 823197-17-7P
823197-18-8P 823197-19-9P 823197-20-2P
823197-21-3P 823197-22-4P 823197-23-5P
823197-24-6P 823197-25-7P 823197-26-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-[(4-substituted piperazine-1-sulfonylmethyl)alkyl]-N-hydroxyformamides as metalloproteinase inhibitors)

RN 823197-00-8 CAPLUS

CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyrimidinyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 823197-01-9 CAPLUS
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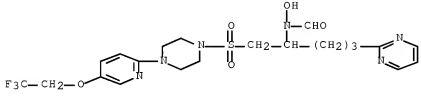
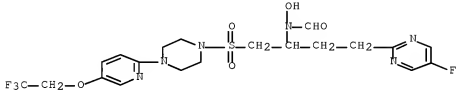
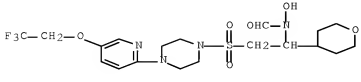


RN 823197-02-0 CAPLUS
CN Piperazine, 1-[[4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyridinyl]- (9CI) (CA INDEX NAME)

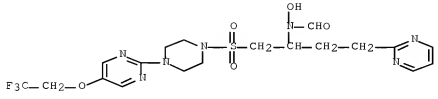


RN 823197-03-1 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-(tetrahydro-2H-pyran-4-yl)ethyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyridinyl]- (9CI) (CA INDEX NAME)

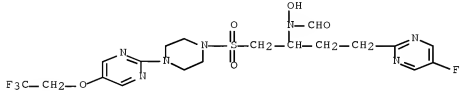


RN 823197-04-2 CAPLUS

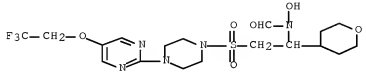
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyrimidinyl]- (9CI) (CA INDEX NAME)



RN 823197-05-3 CAPLUS
CN Piperazine, 1-[[4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyrimidinyl]- (9CI) (CA INDEX NAME)

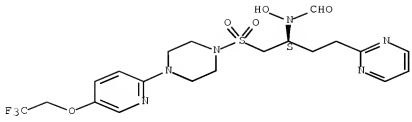


RN 823197-06-4 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-(tetrahydro-2H-pyran-4-yl)ethyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyrimidinyl]- (9CI) (CA INDEX NAME)

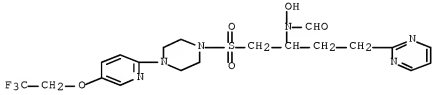


RN 823197-07-5 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyridinyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

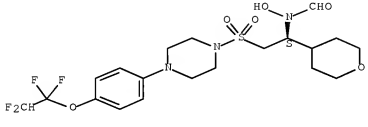


RN 823197-08-6 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyridinyl]- (9CI) (CA INDEX NAME)

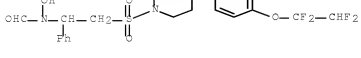


RN 823197-09-7 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-2-(tetrahydro-2H-pyran-4-yl)ethyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

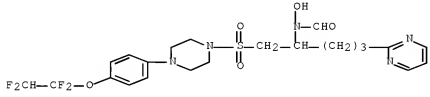
Absolute stereochemistry.



RN 823197-10-0 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-phenylethyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

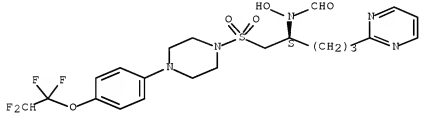


RN 823197-11-1 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

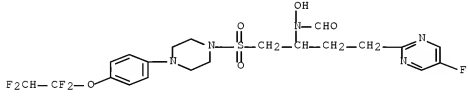


RN 823197-12-2 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

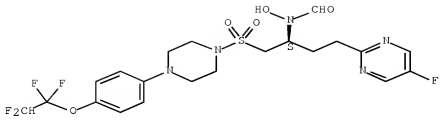


RN 823197-13-3 CAPLUS
CN Piperazine, 1-[[4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

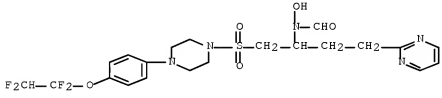


RN 823197-14-4 CAPLUS
CN Piperazine, 1-[[(2S)-4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

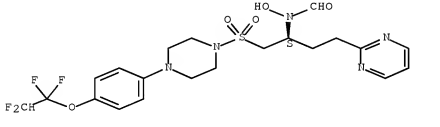


RN 823197-15-5 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

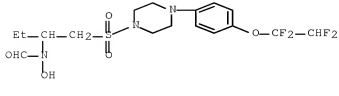


RN 823197-16-6 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

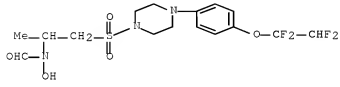
Absolute stereochemistry.



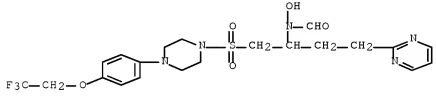
RN 823197-17-7 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)butyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)



RN 823197-18-8 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)propyl]sulfonyl]-4-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

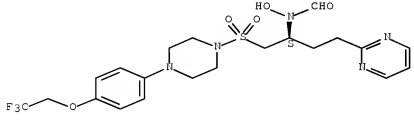


RN 823197-19-9 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

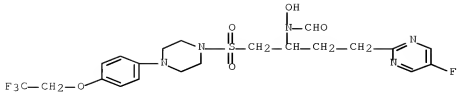


RN 823197-20-2 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-4-(2-pyrimidinyl)butyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

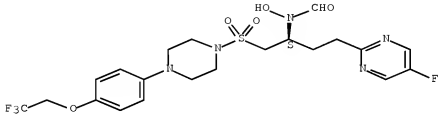


RN 823197-21-3 CAPLUS
CN Piperazine, 1-[[4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

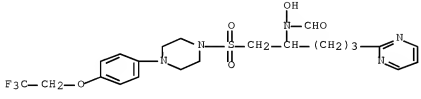


RN 823197-22-4 CAPLUS
CN Piperazine, 1-[[(2S)-4-(5-fluoro-2-pyrimidinyl)-2-(formylhydroxyamino)butyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

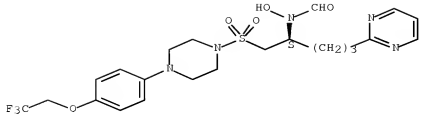


RN 823197-23-5 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)



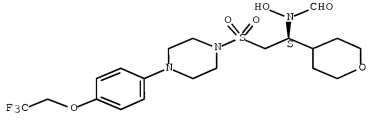
RN 823197-24-6 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

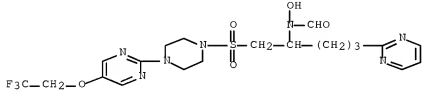


RN 823197-25-7 CAPLUS
CN Piperazine, 1-[[(2S)-2-(formylhydroxyamino)-2-(tetrahydro-2H-pyran-4-yl)ethyl]sulfonyl]-4-[4-(2,2,2-trifluoroethoxy)phenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 823197-26-8 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-[5-(2,2,2-trifluoroethoxy)-2-pyrimidinyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2000:161258 CAPLUS Full-text
DOCUMENT NUMBER: 132:207849
TITLE: Preparation of arylpiperazines as metalloproteinase inhibiting agents (MMP)
INVENTOR(S): Barlaam, Bernard Christophe; Newcombe, Nicholas John; Tucker, Howard; Waterson, David
PATENT ASSIGNEE(S): Zeneca Limited, UK; Zeneca-Pharma Sa
SOURCE: PCT Int. Appl., 82 pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000012478	A1	20000309	WO 1999-GB2801	19990825
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PRIORITY APPLN. INFO.:			EP 1998-402144	A 19980831
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			WO 1999-GB2801	W 19990825
			US 2001-763709	A1 20010226

OTHER SOURCE(S): MARPAT 132:207849
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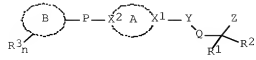

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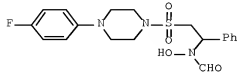


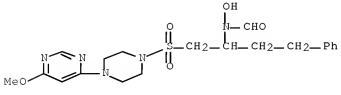
II

AB The title compds. [I; B = monocyclic or bicyclic alkyl, aryl, etc.; R3 = H, halo, NO2. etc.; n = 1-3; P = (CH2)n (wherein n = 0-2), alkene, alkyne, etc.; A = (un)substituted 5-7 membered aliphatic ring; X1, X2 = N, C, where a ring substituent on ring A is a oxo group that is preferably adjacent a ring N atom; Y = SO2, CO; Z = CONHOH, Y = CO and Q = CR6R7, CR6R7CH2, NR6, NR6CH2 (wherein R6 = H, alkyl, aralkyl, etc.; R7 = H, alkyl; R7 together with R6 forms a carbocyclic or heterocyclic spiro 5-7 membered ring, the latter containing at least one heteroatom selected from N, O, S); Z = CONHOH, Y = SO2 and Q = CR6R7, CR6R7CH2; Z = N(OH)CHO and Q = CHR6, CHR6CH2, NR6CH2; R1 = H, alkyl, cycloalkyl, etc.; R2 = H, alkyl, aryl, etc.], useful as metalloproteinase inhibitors (no data), especially as inhibitors of MMP 13, in treating arthritis and atherosclerosis, were prepared E.g., a multi-step synthesis of the title piperazine II was given. Compds. I are effective at 0.5-30 mg/kg/day.

IT 260439-08-5P 260439-96-1P 260440-00-4P
260440-21-9P 260441-00-7P 260441-01-8P
260441-02-9P 260441-03-0P 260441-04-1P
260441-05-2P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of arylpiperazines as metalloproteinase inhibiting agents (MMP))

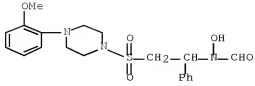
RN 260439-08-5 CAPLUS

CN Piperazine, 1-[[2-(formylhydroxyamino)-4-phenylbutyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)

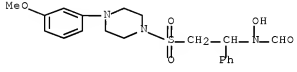


RN 260439-96-1 CAPLUS

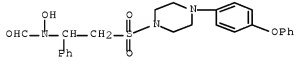
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-phenylethyl]sulfonyl]-4-(2-methoxyphenyl)- (9CI) (CA INDEX NAME)



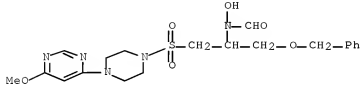
RN 260440-00-4 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-phenylethyl]sulfonyl]-4-(3-methoxyphenyl)- (9CI) (CA INDEX NAME)



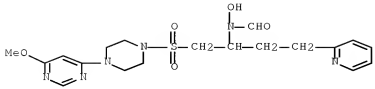
RN 260440-21-9 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-2-phenylethyl]sulfonyl]-4-(4-phenoxyphenyl)- (9CI) (CA INDEX NAME)



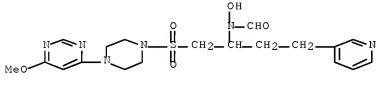
RN 260441-00-7 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-3-(phenylmethoxy)propyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



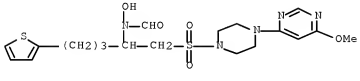
RN 260441-01-8 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(2-pyridinyl)butyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



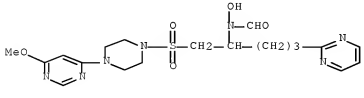
RN 260441-02-9 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-4-(3-pyridinyl)butyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



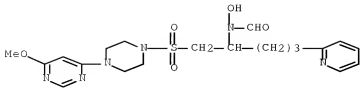
RN 260441-03-0 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-thienyl)pentyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



RN 260441-04-1 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-pyrimidinyl)pentyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



RN 260441-05-2 CAPLUS
CN Piperazine, 1-[[2-(formylhydroxyamino)-5-(2-pyridinyl)pentyl]sulfonyl]-4-(6-methoxy-4-pyrimidinyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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FILE 'REGISTRY' ENTERED AT 18:12:52 ON 15 JAN 2008
L1 STRUCTURE UPLOADED
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